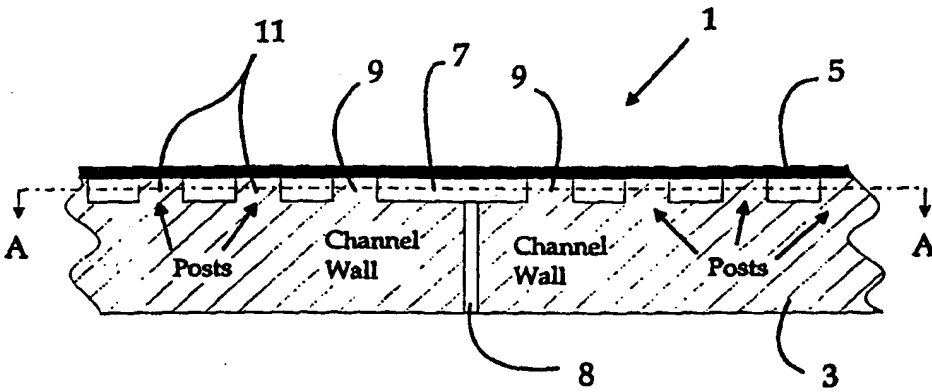




INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

<p>(51) International Patent Classification ⁶ : B01L 3/00, G01N 27/26</p>	<p>A1</p>	<p>(11) International Publication Number: WO 98/32535 (43) International Publication Date: 30 July 1998 (30.07.98)</p>
<p>(21) International Application Number: PCT/SE98/00102 (22) International Filing Date: 23 January 1998 (23.01.98) (30) Priority Data: 9700205-9 24 January 1997 (24.01.97) SE (71)(72) Applicants and Inventors: LINDBERG, Peter [SE/SE]; Serenadvägen 30, S-131 40 NACKA (SE). ROERADE, Johan [SE/SE]; Sågvägen 4, S-147 40 Tumba (SE). STJERNSTRÖM, Mårten [SE/SE]; Styrmansgatan 23, S-114 54 Stockholm (SE). VIOVY, Jean Louis [FR/FR]; Laboratoire de Physico-Chimie, Institut Curie Section de Recherche, 11, rue P & M Curie, F-75231 Paris Cedex 05 (FR). (74) Agent: AWAPATENT AB; P.O. Box 45086, S-104 30 Stockholm (SE).</p>		<p>(81) Designated States: AL, AM, AT, AT (Utility model), AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, CZ (Utility model), DE, DE (Utility model), DK, DK (Utility model), EE, EE (Utility model), ES, FI, FI (Utility model), GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SK (Utility model), SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). Published With international search report.</p>
<p>(54) Title: INTEGRATED MICROFLUIDIC ELEMENT</p>  <p>(57) Abstract</p> <p>An integrated microfluidic element (1) composed of two juxtaposed plates (3, 5) bonded together, wherein at least one plate (3) has an etched structure or pattern of channels (7) on the surface facing the other plate (1) to form sealed micro channels (7), said element comprising micro spacers or posts (11) distributed over the etched surface of said one plate outside of said etched structure or pattern, and walls (9) surrounding said channels (7), said walls (9) having a height equal to that of said spacers or posts (11); and a method for the manufacture of such integrated microfluidic element.</p>		

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece			TR	Turkey
BG	Bulgaria	HU	Hungary	ML	Mali	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MN	Mongolia	UA	Ukraine
BR	Brazil	IL	Israel	MR	Mauritania	UG	Uganda
BY	Belarus	IS	Iceland	MW	Malawi	US	United States of America
CA	Canada	IT	Italy	MX	Mexico	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NE	Niger	VN	Viet Nam
CG	Congo	KE	Kenya	NL	Netherlands	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NO	Norway	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	NZ	New Zealand		
CM	Cameroon			PL	Poland		
CN	China	KR	Republic of Korea	PT	Portugal		
CU	Cuba	KZ	Kazakhstan	RO	Romania		
CZ	Czech Republic	LC	Saint Lucia	RU	Russian Federation		
DE	Germany	LI	Liechtenstein	SD	Sudan		
DK	Denmark	LK	Sri Lanka	SE	Sweden		
EE	Estonia	LR	Liberia	SG	Singapore		

INTEGRATED MICROFLUIDIC ELEMENT

The present invention relates to integrated micro-fluidic elements and a method for the manufacture thereof. Such elements are often composed of two juxtaposed plates bonded together, wherein one plate has an etched structure or pattern of channels on the surface facing the other plate to form sealed microchannels.

Glass substrates have in recent years been used in the manufacture of miniaturized analytic electrophoresis instrumentation with micromachining techniques (Harrison, D. J.; Fluri, K.; Seiler, K.; Fan, Z.; Effenhauser, C. S.; Manz, A., *Science*, **261** (1993) 895-897. Effenhauser, C. S.; Manz, A.; Widmer, H. M., *Anal. Chem.*, **65** (1993) 2637-2642. Seiler, K.; Harrison, D. J.; Manz, A.; *Anal. Chem.*, **65** (1993) 1481-1488. Jacobson, S. C.; Hergenröder, R.; Koutny, L. B.; Ramsey, J. M., *Anal. Chem.*, **66** (1994) 2369-2373. Effenhauser, C. S.; Paulus, A.; Manz, A.; Widmer, H. M., *Anal. Chem.*, **66** (1994) 2949-2953. Seiler, K.; Fan, Z. H.; Fluri, K.; Harrison, D. J., *Anal. Chem.*, **66** (1994) 3485-3491. Jacobson, S. C.; Koutny, L. B.; Hergenröder, R.; Moore, A. W.; Ramsey, J. M., *Anal. Chem.*, **66** (1994) 3472-3476).

The insulating glass material permits the use of high voltages which can accomplish fast and efficient separations and its transparency allows for sensitive on-column optical sample detection.

Bonding of planar structures is an important step necessary in the manufacture of micro-instrumentation. For utilization of microfabricated flow channels in analytical techniques such as electrophoresis, chromatogra-

phy, flow injection analysis or field-flow fractionation, the bond between the structures should preferably be direct. Thus, adhesives should be avoided that can clog the capillaries and negatively effect the efficiency or the liquid flow pattern. Accordingly, an etched structure in for example glass is by preference sealed to another glass plate by fusion bonding at a temperature that permits fusion while not deforming the glass parts, to produce uniformly assembled channel structures.

It is difficult to manufacture large bonded assemblies without irregularities, since the demand on the substrate material in terms of planarity, smoothness and cleanness increases with the area of the substrate. Furthermore, it is usually essential that the surfaces are connected under extreme clean room conditions, so that particle contamination can be eliminated at their interface. Void formation often occurs in the process cycle when bonding starts at the same time at various locations. Once a void is generated the trapped gas cannot be exhausted from its confinement.

The glass material most commonly used in micromachining laboratories is polished substrates of Pyrex Corning 7740, due to its compatibility with silicon in terms of thermal expansion. This expensive material is extensively used for anodic bonding to silicon in the manufacture of microsenors (Cozma, A.; Puers, B. J. *Micromech. Microeng.* 5 (1995) 98-102). However, details on glass-glass fusion bonding of micromachined structures are very sparse in the literature. The reported bonding process is characterized by a low yield which often involves repeated cycles ((Harrison, D. J.; Fluri, K., Seiler, K.; Fan, Z.; Effenhauser, C. S.; Manz, A., *Science*, 261 (1993) 895-897), including the use of weights placed over poorly bonded regions (Fan, Z.; Harrison, D.

J. Anal. Chem **66** (1994) 177-184). Recently, the yield has been improved by the use of sophisticated polishing and cleaning instrumentation (Fluri, K.; Fitzpatrick, G.; Chiem, N.; Harrison, D. J. Anal. Chem **68** (1996) 4285-
5 4290), available only in specialized micromachining laboratories.

For sample detection purposes, many applications require that one of the glass substrates is very thin (0.15 -0.20 mm), e.g. when using high numerical aperture micro-
10 scope objectives. An important example of such an application where a high degree of magnification is required is the direct observation of DNA polymer motion by fluorescence microscopy. Bonding thin glass introduces additional problems mainly due to that the commercially
15 available thin cover glass often is manufactured by a drawing process and therefore not very planar. Raley et al. reported on a etch-back technique where first two thicker sheaths of Corning 7740 were bonded together and subsequently thinning one of the sheaths by etching and
20 several grinding steps (Raley, N. F.; Davidson, J. C.; Balch, J. W. *Proc. SPIE-Int. Soc. Opt. Eng.* **2639** (1995) 40-45). Their best glass-glass bonding results were reported to be in the order of a 85 % area coverage for 5 x 5 cm to 5 x 18 cm glass specimens with a original thickness of 800 μm . However, the etch-back technique is anticipated to depend on how well the etching process can
25 be optimized.

The present invention has for its main object to provide new techniques for the provision of integrated
30 microfluidic elements where the problems encountered with the prior art as illustrated above are eliminated or at least greatly reduced.

Another object of the invention is to provide a method for the manufacture of integrated microfluidic

elements, wherein entrapment of gas between the plates to be bonded together can be avoided.

Yet another object of the invention is to provide a method for such manufacture, wherein the problems encountered in the bonding of two plates of different thermal coefficients of expansion together will be largely avoided.

Still another object of the invention is to provide integrated microfluidic elements free of undesirable voids and less vulnerable to inconsistencies in thermal coefficients of expansion.

For these and other objects which will be clear from the following disclosure the invention provides for a method for the manufacture of an integrated microfluidic element composed of two juxtaposed plates bonded together, wherein at least one plate has an etched structure or pattern of channels on the surface facing the other plate to form sealed microchannels. This method is characterized by forming, distributed over the etched surface of said one plate outside of said etched structure or pattern, micro spacers or posts 11, and by forming walls surrounding said channels 9, said walls 9 having a height equal to that of said spacers or posts 11 and then bonding the two plates 3,5 together to form said element.

The plates can be bonded together by fusion bonding at an increased temperature which does not exceed the softening temperature of the plates.

The plates may also be bonded together by field assisted bonding methods. The bonding techniques used are not critical to the invention and any conventional bonding method can be used. An example of such conventional bonding method is anodic bonding of a glass plate to a silicon substrate.

The plates used can be constituted by materials used in the art, such as ordinary glass, silicon, quartz, dia-

mond, carbon, ceramics or polymers. Particularly useful materials are glass, quartz and silicon.

It is particularly preferred in the method of the invention to form also the spacers or posts and the walls simultaneously with the forming of the structure of pattern of channels by etching.

According to an alternative method the etching can be carried out in two steps, a first step to form the channels and a second step to form the spacers or posts. By such alternative method the depth of the etched sections can be varied.

In some cases it is desired to impart special properties to the channels formed, and here the juxtaposed surfaces of the plates are covered by a thin layer before bonding the plates together. Such layer can be formed e.g. by chemical vaporization deposition (CVD), and the layer can be constituted by any desired material, such as silicon nitride, metals, glass etc.

Access to the channels formed in the microfluidic element of the invention is suitably obtained by the formation of holes in either or both of the two plates in positions coinciding with the channels.

To obtain optimal performance of the microfluidic element of the invention it is preferred that the contact surface between the plates is less than about 50% of the surface of each plate. This surface now referred to is the major side surface of the plate corresponding to the surface of the plate facing the accompanying juxtaposed plate.

The invention also provides for an integrated microfluidic element comprising two juxtaposed plates bonded together, wherein at least one plate has an etched structure or pattern of channels on the surface facing the other plate to form sealed microchannels. Such element is characterized by microspacers or posts distributed over

the etched surface of said one plate outside of said etched structure or pattern, and by walls surrounding said channels. These walls have a height equal to that of said spacers or posts.

5 The posts or "lines" are preferably substantially equally distributed over the etched surface of the plate on areas outside of the etched structure or pattern.

 The present invention as outlined above efficiently reduces or eliminates the problems associated with the
10 prior art techniques. Thus, the use of microspacers or posts distributed over the etched surface of the plate greatly reduces the risk for the formation of voids or cracks in the plate specimens which can be due to dust particles, non-planarity and inconsistencies or differences
15 in thermal coefficients of expansion. Furthermore, the inventive concept results in flexible structures of less built-in tension irrespective of differences in thermal coefficient of expansion of the two plates. Finally, less expensive materials can be used, such as
20 ordinary soda-lime microslide glass which, otherwise, would be virtually impossible to use applying the conventional methods presently available.

 According to a special embodiment of the invention enabling use of a variety of substrates or materials in
25 the plates, the surfaces of the plates facing each other are coated after etching with a thin layer of quartz (SiO_2), such as by chemical vaporization deposition (CVD). In this manner the two plates can be joined together by conventional bonding techniques, such as fusion
30 bonding at an increased temperature. Such method serves the double purpose of enabling easy bonding of the two plates together and simultaneously obtaining a channel lining of a uniform structure. The thickness of such quartz layer may be from fractions of a micron up to about
35 10 microns.

The invention will in the following be further described more in detail by non-limiting examples with reference to the appended drawing, wherein:

Fig. 1 is a section in sideview of an element according to the present invention;

Fig. 2 is a plan view of the element of Fig. 1 taken in a plane along the line A-A in Fig. 1; and

Fig. 3 is the layout of a photomask for a capillary elektrophoretic chip.

While the invention in the following will be exemplified mainly with reference to the use of glass plates for the manufacture of elements in accordance with the present invention it is to be noted that the invention is in no way limited only to the use of such glass plates but is applicable to all types of materials suitable for the intended purpose.

When using glass plates the thickness thereof can vary between about 0.1 and 1 mm, and the deptch of etching can vary between about 1 μ to about 100 μ . The invention greatly facilitates the use of different materials in the two plates, such as one glass plate to be combined with a quartz or silicon plate.

Example 1

The channel manufacturing process includes the steps of: simultaneously HF etching a channel 7 and an array of posts 11 in a glass substrate 3; drilling connection holes 8 to the channel in the glass substrate 3 and bonding the formed channels 7 with a thin cover slip 5 at temperatures below the softening point of the glass (Fig. 1 and 2).

Menzel soda-lime microslide glass plates 3 (76 x 25 mm) with a thickness of 1 mm (\pm 0.1 mm) were used as substrate for etching the channels (7). Cover glass plates (0.17 mm) were also obtained from Menzel. The data given

by the manufacturer for the mean coefficient of thermal expansion are $90.6 \times 10^{-7} \text{ K}^{-1}$ and $(73-74) \times 10^{-7} \text{ K}^{-1}$ for the microslide and cover slips respectively. The corresponding softening points are 720°C and $732-736^\circ\text{C}$.

5 The photo lithographic mask is made with an ordinary CAD program and is printed with a photo setter on a transparent film. The processing chemicals (NH_3 , H_2O_2 , HCl , NH_4F , HF , VLSI Selectipur® grade) were all obtained from Merck (Darmstadt, Germany). All processing solutions
10 were prepared with de-ionized water from a Milli-Q system (Millipore, Bedford, MA) filtered through $0.2 \mu\text{m}$ filters (Millipore).

 The glass substrates were first carefully cleaned in RCA-1 (5 parts distilled H_2O : 1 part NH_3 (25%) : 1 H_2O_2 (20
15 %)) and RCA-2 (6 parts H_2O : 1 part HCl (37%) : 1 part H_2O_2 (20%)) for 10 min respectively and dehydrated in an oven (130°C) for 20 min. For improved adherence of the photoresist, the surfaces were first primed by exposing the substrates with Microposit Primer (Shipley, Marlbor-
20 ough, MA) fumes for 3 min. The glass substrates were then coated with a positive resist (Microposit S1813 Photo Resist, Shipley) with the aid of a lint free paper and softbaked in an oven at (90°C) for 35 min. Next, UV li-
thography followed by a 1 min immersion in a developing
25 bath (Microposit Developer 351, Shipley) and cleaning in distilled H_2O the UV exposed parts of the underlying glass were laid open for subsequent etching. After a hardbaking step (130°C) for 45 min, etching was performed in a vigorously stirred aqueous mixture of 5% buffered HF
30 (7 parts 40 % NH_4F ; 1 part 50% HF) and 9.25% conc. HCl for 45 - 60 min at room temperature to form channels 7, walls 9 and posts 11. The resulting etch depth was approximately $80 \mu\text{m}$.

Connection holes 8 to the flow channel were manufactured by drilling the microslide with a carbide-drill steel (0.5 mm diameter). The remaining drilling dust was removed with ultrasonic, RCA-1 and RCA-2 rinsing steps.

5 Bonding was performed by first carefully mating the wafers in a clean hood and placing the substrates with the cover glass downwards in an oven at 630 ° C for 8 hours. In order to avoid fusion to and replication of scratches from the underlying support, plates of polished vitreous

10 carbon were utilized (V25 grade, Le Carbone-Lorraine, Gennevilliers, France).

Example 2

The use of the new bonding technique to manufacture microfluidic structures is illustrated in this example.

15 The glass channel is designed for directly observing fluorescent images of individual DNA polymers undergoing separations, under a high numerical aperture microscope. The photomask used to manufacture this capillary electrophoretic chip is shown in Fig. 3 as a layout of the photomask for the capillary electrophoretic chip. The dark

20 areas define the channel walls 9 and the posts 11 and the channel dimensions are 500 μm x 3 mm for the large channels 17 and 50 μm x 40 mm for the thin channel. The buffer and sample reservoirs 21,23, respectively, have a

25 diameter of 2 mm. The square posts dimensions are 400 μm x 400 μm .

The photolithographic technique presented here does not involve any advanced deposition or mask-alignment steps. The process is facilitated by using a positive resist as a direct etch mask instead of using the commonly

30 used chromium/gold coating. The pattern in the photomask is simply transferred by UV lithography to a film of photosensitive positive resist which, in turn, by the etching process conveys the geometric shape to the glass sub-

strate. The positive resist withstands diluted buffered HF etch solution at least up to 1 hour in room temperature, when concentrated HCl is added to the solution. This etch time is sufficient to produce deep etched channel structures in ordinary sodalime microscope slides. We observed that the lower pH also was beneficial for the etching process itself. At the used concentration of buffered HF, the smoothness of the channel walls was significantly improved in comparison with the case where no HCl addition was made. Additionally, at higher pH and HF concentrations we observed crystalline precipitates which was determined to be CaF_2 with EDAX.

No remaining voids or breakage after the thermal bonding step were observed when the array of posts was utilized as underpinning elements. It is suggested that the main reason for this observation is due to the fact that gases are permitted to be exhausted through the two-dimensional network of open channels. Thus, gas expansion in enclosed voids never appears when the substrates are heated. Additionally, no cracks were introduced during the bonding step due to mismatch in the thermal expansion coefficients between the etched substrate and the cover glass material. This problem is also diminished by the lattice of raised posts, since their higher degree of flexibility reduces the strain imposed on the interface between the substrates. This opens the possibility to fusion bond materials with larger difference in thermal expansion coefficients such as quartz - glass, silicon - glass etc. and broadens the choice of glass material for the anodic bonding technique (Cozma, A.; Puers, B. J. *Microeng. 5* (1995) 98-102), which also is performed at elevated temperatures.

It is to be observed that the present invention is not restricted to the specific embodiments described

11

above but is broadly applicable, and the invention is not to be construed to be limited otherwise than specified in the appended patent claims.

CLAIMS

1. A method for the manufacture of an integrated microfluidic element (5) composed of two juxtaposed plates (3,5) bonded together, wherein at least one plate (3) has an etched structure or pattern of channels (7) on the surface facing the other plate (1) to form sealed micro channels (7), characterized by forming, distributed over the etched surface of said one plate outside of said etched structure or pattern, micro spacers or posts (11), and by forming walls (9) surrounding said channels (7), said walls (9) having a height equal to that of said spacers or posts (11) and then bonding the two plates (3,5) together to form said element.
2. A method according to claim 1, wherein the plates (3,5) are bonded together by fusion bonding at an increased temperature not exceeding the softening temperature of the plates.
3. A method according to claim 1 or 2, wherein the plates (3,5) are bonded together by field-assisted bonding techniques.
4. A method according to any preceding claim, wherein the plates (3,5) are constituted by materials selected from ordinary glass, quartz and silicon.
5. A method according to any preceding claim, wherein also said spacers or posts (11) and walls (9) are formed by etching.
6. A method according to claim 5, wherein the etching is carried out in one step to form simultaneously both channels (7) and spacers or posts (11).
7. A method according to claim 5, wherein the etching is carried out in two steps, a first step to form the channels (7) and a second step to form the spacers or posts (11).
8. A method according to any preceding claim, wherein the juxtaposed surfaces of the plates (3,5) before

bonding the plates together are covered by a thin layer to form a channel lining of the desired properties.

9. A method according to claim 8, wherein said thin layer is formed by chemical vaporization deposition
5 (CVD).

10. A method according to any preceding claim, wherein excess to the channels formed is obtained by forming holes (8) in either or both of the two plates (3,5).

11. A method according to any preceding claim, wherein the contact surface between the plates (3,5) is less
10 than about 50% of the surface of each plate.

12. An integrated microfluidic element (1) composed of two juxtaposed plates (3,5) bonded together, wherein at least one plate (3) has an etched structure or pattern
15 of channels (7) on the surface facing the other plate (5) to form sealed micro channels (7), characterized by micro spacers or posts (11) distributed over the etched surface of said one plate outside of said etched structure or pattern, and by walls (9) surrounding said channels (7),
20 said walls (9) having a height equal to that of said spacers or posts (11).

13. An integrated microfluid element according to claim 12, wherein the plates (3,5) are constituted by materials selected from ordinary glass, quartz or silicon.

25 14. An integrated microfluid element according to claim 12 or 13, wherein the juxtaposed surfaces of the plates (3,5) before bonding the plates together have been covered by a thin layer to give a channel lining of the desired properties.

30 15. An integrated microfluid element according to any one of claims 12 to 14, wherein excess to the channels is obtained by holes (8) made in either or both of the two plates (3,5).

35 16. An integrated microfluid element according to any one of claims 12 to 15, wherein the contact surface between the plates (3,5) is less than about 50% of the surface of each plate.

1/1

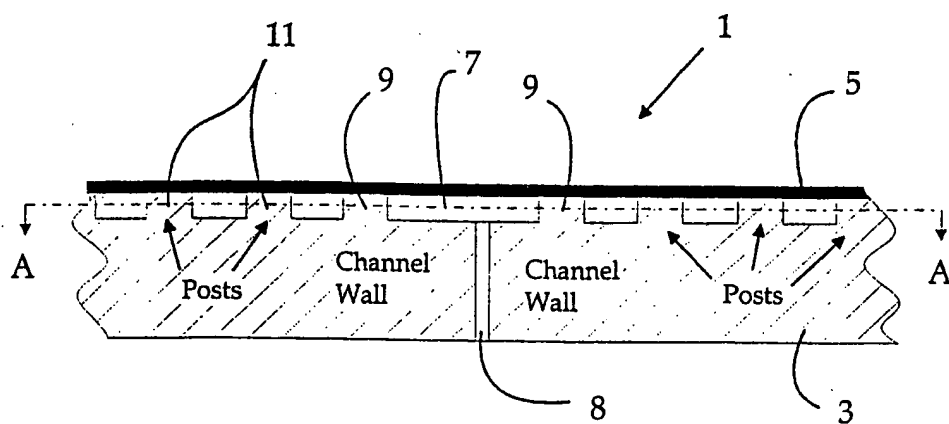


FIG. 1

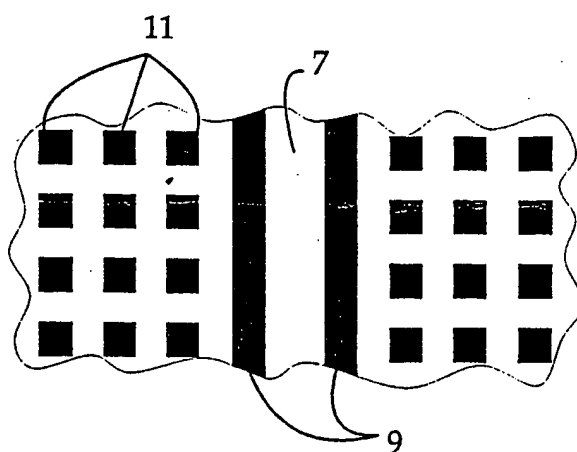


FIG. 2

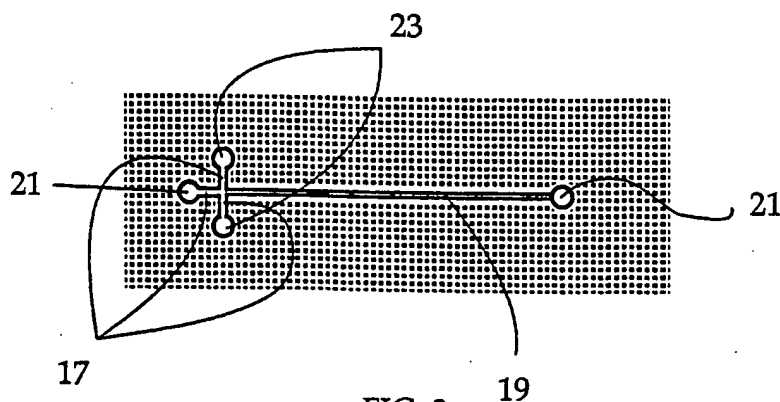


FIG. 3

INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 98/00102

A. CLASSIFICATION OF SUBJECT MATTER

IPC6: B01L 3/00, G01N 27/26

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC6: B01L, G01N

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

SE,DK,FI,NO classes as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

WPI, EPDOC

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 9116966 A1 (PHARMACIA BIOSENSOR AB), 14 November 1991 (14.11.91) --	1,12
A	Analusis, Volume 21, 1993, RH Austin et al, "Electrophoresis and microlithography" page 236, column 1, line 1 - column 1, line 64; page 236, column 2, line 1 - column 2, line 28 --	1-16
A	US 5593838 A (PETER J.ZANZUCCHI ET AL), 14 January 1997 (14.01.97) --	1-16
A	WO 9322055 A2 (TRUSTEES OF THE UNIVERSITY OF PENNSYLVANIA), 11 November 1993 (11.11.93) --	1-16

☐ Further documents are listed in the continuation of Box C.
 ☒ See patent family annex.

* Special categories of cited documents:	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"A" document defining the general state of the art which is not considered to be of particular relevance	"X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"E" earlier document but published on or after the international filing date	"Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"&" document member of the same patent family
"O" document referring to an oral disclosure, use, exhibition or other means	
"P" document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search	Date of mailing of the international search report
15 April 1998	04-05-1998
Name and mailing address of the ISA/ Swedish Patent Office Box 5055, S-102 42 STOCKHOLM Facsimile No. +46 8 666 02 86	Authorized officer Ulf Nyström Telephone No. +46 8 782 25 00

INTERNATIONAL SEARCH REPORT
Information on patent family members

02/04/98

International application No.

PCT/SE 98/00102

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9116966 A1	14/11/91	AT 130528 T DE 69114838 D,T EP 0527905 A,B SE 470347 B,C SE 9001699 A US 5376252 A	15/12/95 05/06/96 24/02/93 31/01/94 11/11/91 27/12/94
US 5593838 A	14/01/97	AU 4152396 A AU 4233796 A CA 2204912 A CA 2205066 A EP 0791238 A EP 0808456 A US 5585069 A US 5643738 A US 5681484 A WO 9615450 A WO 9615576 A	06/06/96 06/06/96 23/05/96 23/05/96 27/08/97 26/11/97 17/12/96 01/07/97 28/10/97 23/05/96 23/05/96

INTERNATIONAL SEARCH REPORT
Information on patent family members

02/04/98

International application No.

PCT/SE 98/00102

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9322055 A2	11/11/93	AT 140025 T	15/07/96
		AT 140880 T	15/08/96
		AT 155711 T	15/08/97
		AU 674685 B	09/01/97
		AU 677197 B	17/04/97
		AU 677780 B	08/05/97
		AU 677781 B	08/05/97
		AU 680195 B	24/07/97
		AU 4222393 A	29/11/93
		AU 4222593 A	29/11/93
		AU 4222693 A	29/11/93
		AU 4222793 A	29/11/93
		AU 4223593 A	29/11/93
		CA 2134474 A	11/11/93
		CA 2134475 A	11/11/93
		CA 2134476 A	11/11/93
		CA 2134478 A	11/11/93
		DE 69303483 D,T	06/02/97
		DE 69303898 D,T	20/02/97
		DE 69312483 D,T	12/02/98
		EP 0637996 A,B	15/02/95
		EP 0637997 A	15/02/95
		EP 0637998 A,B	15/02/95
		EP 0637999 A	15/02/95
		EP 0639223 A,B	22/02/95
		HK 16897 A	13/02/97
		JP 7506256 T	13/07/95
		JP 7506257 T	13/07/95
		JP 7506258 T	13/07/95
		JP 7506430 T	13/07/95
		JP 7506431 T	13/07/95
		US 5304487 A	19/04/94
		US 5635358 A	03/06/97
		US 5726026 A	10/03/98
		WO 9322053 A	11/11/93
		WO 9322054 A	11/11/93
		WO 9322058 A	11/11/93
		WO 9322421 A	11/11/93
		US 5296375 A	22/03/94
		US 5427946 A	27/06/95
		US 5498392 A	12/03/96
		US 5587128 A	24/12/96
		US 5486335 A	23/01/96
		US 5637469 A	10/06/97